CLAIMS

1. A compound of formula (I):

10 (I)

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in which:

X is N, NH, :CH or CH₂;

Y is N, :CH, CO, CH₂ or :CNR²R³, where R² and R³ are independently hydrogen, C_{1-6} alkyl or C_{3-6} cycloalkyl;

R is aryl or heteroaryl optionally substituted by halogen, amino, hydroxy, cyano, nitro, trifluoromethyl, carboxy, CONR 5 R 6 , SO $_2$ NR 5 R 6 , SO $_2$ R 4 , NHSO $_2$ R 4 , NHCOR 4 , ethylenedioxy, methylenedioxy, C $_{1-6}$ alkyl, C $_{1-6}$ alkoxy, SR 4 or NR 5 R 6 where R4 is hydrogen, C $_{1-6}$ alkyl or C $_{3-6}$ cycloalkyl, R 5 and R 6 are independently hydrogen, C $_{1-6}$ alkyl or together with the nitrogen atom to which they are attached form a 5- or 6-membered saturated ring optionally containing a further O, S or NR 4 group; or R is hydrogen, C $_{1-6}$ alkyl or C $_{3-6}$ cycloalkyl both of which can optionally contain one or more O, S or NR 4 groups,

 R^1 is a group Y(CH₂)pR⁷ where p is 0, 1 or 2 and Y is O or NR⁸ where R⁸ is hydrogen, C₁₋₆ alkyl or C₃₋₆ cycloalkyl;

and R⁷ is a 5- or 6-membered saturated ring containing one or more O, S or N atoms, aryl or a heteroaryl group containing one to four heteroatoms selected from O, S or N, the saturated ring, aryl and heteroaryl groups all being optionally substituted by halogen,

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amino, hydroxy, cyano, nitro, trifluoromethyl, carboxy, CONR⁵R⁶, SO₂NR⁵R⁶, SO₂R⁴, NHSO₂R⁴, NHCOR⁴, C₁₋₆ alkyl, C₁₋₆ alkoxy, SR⁴ or NR⁵R⁶ where R4 is hydrogen, C₁₋₆ alkyl or C₃₋₆ cycloalkyl, R⁵ and R⁶ are independently hydrogen, C₁₋₆ alkyl or together with the nitrogen atom to which they are attached form a 5- or 6-membered saturated ring optionally containing a further O, S or NR⁴ group;

or R¹ is a group NR⁹R¹⁰ where R⁹ and R¹⁰ are independently hydrogen or C₁₋₆ alkyl optionally containing one or more O, S or NR⁴ groups, or R⁹ and R¹⁰ together with the nitrogen atom to which they are attached form a 5 or 6-membered saturated ring optionally containing a further O, S or N atom and optionally substituted by NR⁹R¹⁰, CO₂C₁₋₆ alkyl, CONR¹¹R¹² where R¹¹ and R¹² are independently hydrogen or C₁₋₆ alkyl, aryl or heteroaryl group optionally substituted by halogen, amino, hydroxy, cyano, nitro, trifluoromethyl, carboxy, CONR⁵R⁶, SO₂NR⁵R⁶, SO₂R⁴, NHSO₂R⁴, NHCOR⁴, C₁₋₆ alkyl, C₁₋₆ alkoxy, SR⁴ or NR⁵R⁶ where R4 is hydrogen, C₁₋₆ alkyl or C₃₋₆ cycloalkyl, R⁵ and R⁶ are independently hydrogen, C₁₋₆ alkyl or together with the nitrogen atom to which they are attached form a 5- or 6-membered saturated ring optionally containing a further O, S or NR⁴ group; and pharmaceutically acceptable salts or solvates thereof.

- 2. A compound according to claim 1 in which X is N and Y is :CH, X and Y are:CH or X and Y are CH₂
 - 3. A compound according to claim 1 or 2 in which R is C₁₋₄alkyl, or phenyl substituted by halogen, SO₂Me, C₁₋₆alkoxy or C₁₋₄alkyl.
- 4. A compound according to any one of claims 1 to 3 in which R¹ is a group Y(CH₂)pR⁷ where p is 0 and Y is NR⁸ where R⁸ is hydrogen and R⁷ is substituted phenyl.
 - 5. A compound according to any one of claims 1 to 3 in which R¹ is NR⁹R¹⁰ where R⁹ and R¹⁰ are hydrogen or C₁₋₃ alkyl or together with the nitrogen atom to which they are attached form a 5 or 6-membered saturated ring optionally containing a O, S or NR⁴.
 - 6. A compound of formula (I) selected from:
 - 1-[9-(4-Chlorophenyl)-2-cyano-9H-purin-6-yl]-L-prolinamide,
 - 9-(4-Chlorophenyl)-6-(4-pyrrolidin-1-ylpiperidin-1-yl)-9H-purine-2-carbonitrile.
- 9-(4-Chlorophenyl)-6-[(3-pyrrolidin-1-ylpropyl)amino]-9H-purine-2-carbonitrile, 6-(4-Aminopiperidin-1-yl)-9-(4-chlorophenyl)-9H-purine-2-carbonitrile,

- 6-[(2-Aminoethyl)amino]-9-(4-chlorophenyl)-9H-purine-2-carbonitrile,
- 9-(4-Chlorophenyl)-6-(dimethylamino)-9H-purine-2-carbonitrile,
- 9-(4-Methylphenyl)-6-pyrrolidin-1-yl-9H-purine-2-carbonitrile,
- 9-(4-Methoxyphenyl)-6-pyrrolidin-1-yl-9H-purine-2-carbonitrile,
- 9-(4-chlorophenyl)-6-pyrrolidin-1-yl-9H-purine-2-carbonitrile,
 - 9-(4-Chlorophenyl)-6-(ethylamino)-9H-purine-2-carbonitrile,
 - tert-Butyl 4-[9-(4-chlorophenyl)-2-cyano-9H-purin-6-yl]piperazine-1-carboxylate,
 - 9-(4-Chlorophenyl)-6-piperazin-1-yl-9H-purine-2-carbonitrile,
 - 9-(2-Chlorophenyl)-6-morpholin-4-yl-9H-purine-2-carbonitrile
- 9-(3,4-Difluorophenyl)-6-morpholin-4-yl-9H-purine-2-carbonitrile,
 - 9-(4-Isopropylphenyl)-6-morpholin-4-yl-9H-purine-2-carbonitrile,
 - 9-(4-Methoxyphenyl)-6-morpholin-4-yl-9H-purine-2-carbonitrile,
 - 9-(3-Chlorophenyl)-6-morpholin-4-yl-9H-purine-2-carbonitrile,
 - 9-[4-(Methylsulfonyl)phenyl]-6-morpholin-4-yl-9H-purine-2-carbonitrile,
- 5 6-[(4-Chlorophenyl)amino]-9-ethyl-9H-purine-2-carbonitrile.
 - 9-(4-Chlorophenyl)-6-morpholin-4-yl-9H-purine-2-carbonitrile,
 - 8-Amino-6-[(4-chlorophenyl)amino]-9-ethyl-9H-purine-2-carbonitrile.
 - 8-Amino-9-(4-chlorophenyl)-6-morpholin-4-yl-9H-purine-2-carbonitrile,
 - 9-(4-Chlorophenyl)-6-morpholin-4-yl-8-oxo-8,9-dihydro-7H-purine-2-carbonitrile,
- 9-(4-Chlorophenyl)-8-(dimethylamino)-6-morpholin-4-yl-9H-purine-2-carbonitrile,
 - 7-(4-Chlorophenyl)-4-morpholin-4-yl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile,
 - 7-(4-Chlorophenyl)-4-(ethylamino)-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile,
 - 4-[(4-Chlorophenyl)amino]-7-ethyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile.
 - 1-[7-(4-Chlorophenyl)-2-cyano-6,7-dihydro-5H-pyrrolo[2,3-d]pyrimidin-4-yl]-L-
- 25 prolinamide,
 - 1-[2-Cyano-7-(4-methoxyphenyl)-6,7-dihydro-5H-pyrrolo[2,3-d]pyrimidin-4-yl]-L-prolinamide,
 - 7-(4-Methoxyphenyl)-4-pyrrolidin-1-yl-6,7-dihydro-5H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile,
- 7-(4-Methoxyphenyl)-4-morpholin-4-yl-6,7-dihydro-5H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile,
 - 1-(4-Methylphenyl)-4-morpholin-4-yl-1H-pyrazolo[3,4-d]pyrimidine-6-carbonitrile, and pharmaceutically acceptable salts thereof.
- 7. A compound of formula (I) as defined in any one of claims 1 to 6 for use in therapy.

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- 8. A compound of formula (I) as defined in any one of claims 1 to 6 for use in the treatment of pain.
- 9. A compound of formula (I) as defined in any one of claims 1 to 6 for use in the treatment of neuropathic pain.

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- 10. A pharmaceutical composition which comprises a compound of the formula (I) as defined in any one of claims 1 to 6 or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable diluent or carrier.
- 11. A method for producing inhibition of a cysteine protease in a mammal, such as man, in need of such treatment, which comprises administering to said mammal an effective amount of a compound as defined in any one of claims 1 to 6, or a pharmaceutically acceptable salt thereof.
- 12. A method for treating pain, such as neuropathic pain, in a mammal, such as man, in need of such treatment, which comprises administering to said mammal an effective amount of a compound as defined in any one of claims 1 to 6, or a pharmaceutically acceptable salt thereof.
- 13. Use of a compound of the formula (I) as defined in any one of claims 1 to 6 or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for use in the inhibition of Cathepsin S in a warm blooded animal, such as man.